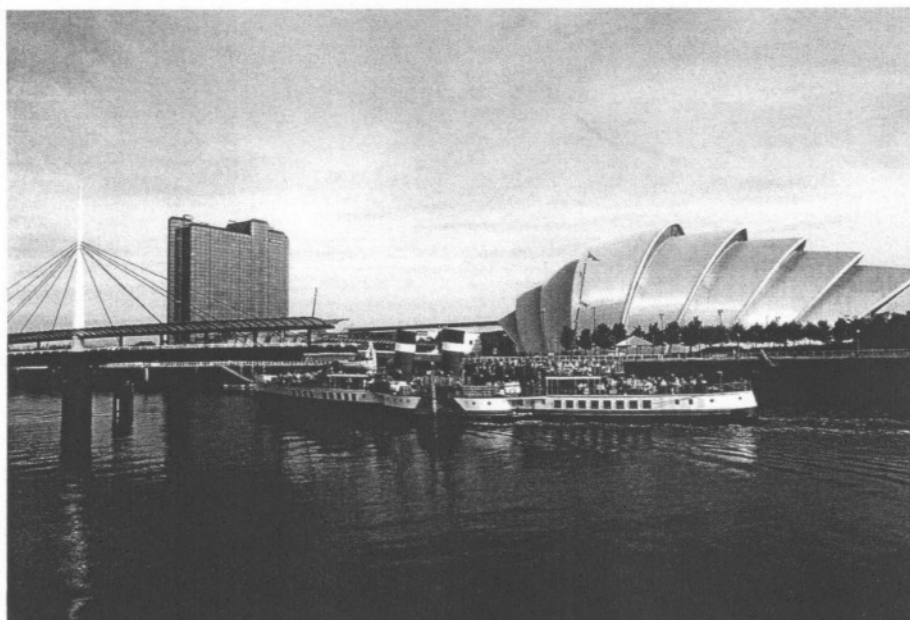


23rd International Congress on  
Heterocyclic Chemistry  
GLASGOW, SCOTLAND

**ICHHC2011**

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*31st July – 4th August 2011*



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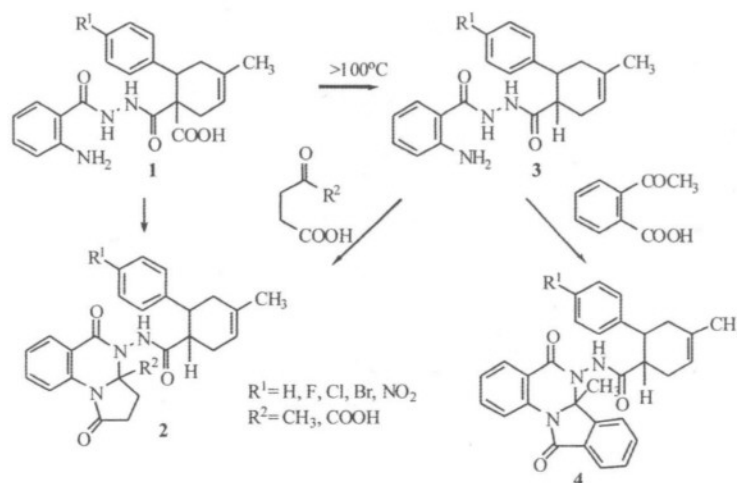
**Programme & Abstract Book**

### Synthesis and Reactivity of N'-acylanthranylhydrazides

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Polyheterocyclic structures containing fused quinazolines are attractive molecular scaffolds due to their broad spectrum of biological activities. Thus, pyrrolo-quinazoline fragment can be found in natural alkaloids<sup>1</sup> and in pharmaceutically active substances.<sup>2</sup> This has led to extensive studies of their synthesis. We proceeded to the synthesis of novel derivatives of 4-amino-tetrahydro-pyrrolo[1,2-*a*]quinazolines which can be viewed as amide-linked-conjugates between the above mentioned fused heterocyclic system and a carbocycle. Starting from commercially available Meldrum's acid and benzaldehydes, 2'-N-substituted anthranylhydrazides **1** can be synthesised in 4 steps.<sup>3</sup> The desired cyclohexenecarbonylamino-tetrahydro-pyrrolo[1,2-*a*]quinazolines **2** can be prepared with aliphatic keto acids in final step.



We have found that compounds **2**, prepared from **1**, are in low yields. Hence, the further experiments were carried out after a decarboxylation of hydrazides **1** in pyridine to give the corresponding 4-amino-tetrahydro-pyrrolo[1,2-*a*]quinazolines **2** in high yields. Extension of this reaction with 2-acetylbenzoic acid gave the related tetrahydroisindolo[2,1-*a*]quinazolin-2(1H)-ones **4**.

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